The listing of claims will replace all prior versions, and listings, of claims in the application:

## **Listing of Claims:**

1. (Previously Presented) A process for preparing tetrahydropterin of the following formula

or a tetrahydropterin compound of said tetrahydropterin that is substituted at the 6-, or 7- or 6and 7- position or positions,

comprising hydrogenating pterin of the following formula

$$\begin{array}{c|c}
H & N & 5 \\
N & 3 & N & 6 \\
H_2 & N & N & 8
\end{array}$$

or a pterin compound of said pterin that is monosubstituted at the 6-, or 7- or 6- and 7- position or positions,

with hydrogen in a polar reaction medium in the presence of a hydrogenation catalyst that is a metal complex that is soluble in the reaction medium, wherein the catalyst contains a ligand which is (i) triarylphosphine, (ii) tetramethylene phenylphosphine (iii) pentamethylene phenylphosphine, or (iv) a bidentate ligand with a tertiary amine group and a phosphine group or with two tertiary phosphine groups as complexing groups, wherein the bidentate ligands form together with a metal atom a five- to ten membered ring.

- 2. (Previously Presented) A process according to claim 1, wherein the polar reaction medium is an aqueous or alcoholic reaction medium.
- 3. (Previously Presented) A process according to claim 1, wherein the pterin compound is folic acid, a folic acid salt, a folic acid ester, a folic acid ester salt or a dihydro form thereof, with the proviso that in the event of using folic acid, a carboxylic acid thereof or a dihydro form thereof, the reaction medium is aqueous, and in the event of using a folic acid ester, a folic acid ester salt or a dihydro form thereof, the reaction medium is an alcohol.
- 4. (Previously Presented) A process according to claim 1, wherein the metal complex contains a chiral ligand.
- 5. (Previously Presented) A process according to claim 3, wherein the metal complex contains a chiral ligand.
- 6. (Previously Presented) A process according to claim 5, wherein the folic acid ester salt is of formula III and is in the form of a single enantiomer or a mixture of enantiomers of formula III,

in which

one of  $R_1$  or  $R_2$  is H, and the other one of  $R_1$  or  $R_2$  is a monovalent hydrocarbon radical or a hydrocarbon radical attached via a carbon atom in which one or more carbon atoms are each independently replaced by oxygen, sulfur, NH, -N=, or -N( $C_1$ - $C_4$  Alkyl)-, or both  $R_1$  and  $R_2$  independently of one another represent a monovalent hydrocarbon radical or a hydrocarbon radical attached via a carbon atom in which one or more carbon atoms are each independently replaced by oxygen, sulfur, NH, -N=, or -N( $C_1$ - $C_4$  Alkyl)-, HA stands for a monobasic to tribasic inorganic or organic acid, and x denotes an integer from 1 to 6 or a fractional number between 0 and 6.

- 7. (Previously Presented) A process according to claim 6, wherein HA is unsubstituted or substituted phenylsulphonic acid.
- 8. (Previously Presented) A process according to claim 1, wherein said process is carried out at a hydrogen pressure of 1 to 500 bars.
- 9. (Previously Presented) A process according to claim 1, wherein said process is carried out at a temperature is 0 to  $150^{\circ}$  C.

- 10. (Previously Presented) A process according to claim 1, wherein the molar ratio of pterin or pterin compound to catalyst is 10 to 100,000.
- 11. (Previously Presented) A process according to claim 1, wherein the reaction medium is water or water in admixture with an organic solvent.
- 12. (Previously Presented) A process according to claim 2, wherein the alcoholic reaction medium is an alcohol, or an alcohol in admixture with an organic solvent.
- 13. (Previously Presented) A process according to claim 1, wherein the metal complex contains a d-8 metal.
  - 14-28. (Cancelled)
- 29. (Previously Presented) A process for preparing tetrahydropterin of the following formula

$$\begin{array}{c|c}
H & H \\
N_3 & N_5
\end{array}$$

$$\begin{array}{c|c}
H & N_8 \\
N_8 & N_8
\end{array}$$

or a tetrahydropterin compound of said tetrahydropterin that is substituted at the 6-, or 7- or 6and 7- position or positions,

comprising hydrogenating pterin of the following formula

or a pterin compound of said pterin that is monosubstituted at the 6-, or 7- or 6- and 7- position or positions,

with hydrogen in alcohol or in alcohol in admixture with an organic solvent in the presence of a hydrogenation catalyst that is a metal complex that is soluble in the reaction medium.

30-32. (Cancelled)

- 33. (Previously Presented) A process according to claim 3, wherein the hydrogenation is carried out at elevated pressure.
- 34. (Previously Presented) A process according to claim 1, wherein the metal complex contains iridium, rhodium or ruthenium.

35-36. (Cancelled)

37-39. (Cancelled)

40. (Previously Presented) A process for preparing tetrahydropterin of the following formula

$$\begin{array}{c|c}
H & H \\
N & 3 \\
H_2 & N \\
1 & H
\end{array}$$

or a tetrahydropterin compound of said tetrahydropterin that is substituted at the 6-, or 7- or 6- and 7- position or positions,

comprising hydrogenating pterin of the following formula

or a pterin compound of said pterin that is monosubstituted at the 6-, or 7- or 6- and 7- position or positions,

with hydrogen in alcohol or in alcohol in admixture with an organic solvent in the presence of a hydrogenation catalyst that is a metal complex that is soluble in the reaction medium, wherein the pterin compound is folic acid, a folic acid salt, a folic acid ester, a folic acid ester salt or a dihydro form thereof, with the proviso that in the event of using folic acid, a carboxylic acid thereof or a dihydro form thereof, the reaction medium is aqueous, and in the event of using a folic acid ester, a folic acid ester salt or a dihydro form thereof, the reaction medium is an alcohol.

## 41-44. (Cancelled)

45. (Previously Presented) A process according to claim 1, wherein the pterin compound is a pterin that is substituted in the 6- position.

46. (Previously Presented) A process according to claim 1, wherein the pterin compound is of formula (A)

$$\begin{array}{c|c}
H & N & R_{100} \\
H_2N & N & R_{101}
\end{array}$$
(A)

in which

 $R_{101}$  is H or independently has the meaning of  $R_{100}$ , and

R<sub>100</sub> is an organic radical attached via a C, O or N atom and having 1 to 50 carbon atoms.

47. (Previously Presented) A process according to claim 46, wherein  $R_{100}$  contains 1 to 30 carbon atoms and is not interrupted or is interrupted by one or more of -O-, -NH-, -N(C<sub>1</sub>-C<sub>4</sub>-alkyl)-, -C(O)-, -C(O)O-, -OC(O)-, -OC(O)O-, -C(O)NH-, -NHC(O)-, -NHC(O)O-, -OC(O)NH-, -NHC(O)NH-, -C(O)N(C<sub>1</sub>-C<sub>4</sub>-alkyl)-, -N(C<sub>1</sub>-C<sub>4</sub>-alkyl)C(O)-, -N(C<sub>1</sub>-C<sub>4</sub>-alkyl)C(O)O-, -OC(O)N(C<sub>1</sub>-C<sub>4</sub>-alkyl)-, -N(C<sub>1</sub>-C<sub>4</sub>-alkyl)C(O)N(C<sub>1</sub>-C<sub>4</sub>-alkyl)-, and which is unsubstituted or is substituted with F, Cl, Br, -CN, -OCN, -NCO, -OH, -NH<sub>2</sub>, -NHC<sub>1</sub>-C<sub>4</sub>-alkyl, -N(C<sub>1</sub>-C<sub>4</sub>-alkyl)<sub>2</sub>, C<sub>1</sub>-C<sub>4</sub>-alkyl, C<sub>1</sub>-C<sub>4</sub>-haloalkyl, C<sub>1</sub>-C<sub>4</sub>-hydroxyalkyl, C<sub>1</sub>-C<sub>4</sub>-alkoxy, C<sub>1</sub>-C<sub>4</sub>-haloalkoxy, -C(O)OH, -C(O)OM<sub>100</sub>, -C(O)OC<sub>1</sub>-C<sub>4</sub>-alkyl, -C(O)NH<sub>2</sub>, -C(O)NHC<sub>1</sub>-C<sub>4</sub>-alkyl, -C(O)N(C<sub>1</sub>-C<sub>4</sub>-alkyl)<sub>2</sub>, R<sub>102</sub>-C(O)O-, R<sub>102</sub>-OC(O)O-, R<sub>102</sub>-C(O)NH-, R<sub>102</sub>-C(O)N(C<sub>1</sub>-C<sub>4</sub>-alkyl)-, R<sub>102</sub>-NHC(O)NH-, R<sub>103</sub>C(O)- or -CH(O), wherein

M<sub>100</sub> is Li, K, Na, NH<sub>4</sub><sup>+</sup>, or ammonium with 1 to 16 carbon atoms,

 $R_{102}$  is  $C_1$ - $C_8$ -alkyl,  $C_5$ - or  $C_6$ -cycloalkyl, phenyl or benzyl, and

 $R_{103}$  is  $C_1$ - $C_4$ -alkyl, phenyl or benzyl.

48. (Previously Presented) A process for preparing tetrahydropterin of the following formula

or a tetrahydropterin compound of said tetrahydropterin that is substituted at the 6-, or 7- or 6- and 7- position or positions,

comprising hydrogenating pterin of the following formula

or a pterin compound of said pterin that is monosubstituted at the 6-, or 7- or 6- and 7- position or positions,

with hydrogen in a polar reaction medium in the presence of a hydrogenation catalyst that is a metal complex that is soluble in the reaction medium of formula XLIV, XLIVa or XLIVb,

$$[X_7Me_2YZ]$$
 (XLIV),  $[X_7Me_2Y]^+A_2^-$  (XLIVa)  $[X_7Ru(II)X_8X_9](XLIVb)$ ,

in which

Y stands for monoolefin ligands or a diene ligand;

 $X_7$  represents an achiral or chiral ditertiary diphosphine, that forms a 5 to 7 membered ring with the metal atom Me<sub>2</sub> or Ru;

 $X_7$  represents an achiral or chiral ligand that forms a 5 to 7 membered ring with the metal atom Me<sub>2</sub> or Ru, wherein said ligand contains two tertiary phosphine groups;

 $Me_2$  denotes Ir(I) or Rh(I);

Z represents -Cl, -Br, or -I; and

 $A_2$  is  $ClO_4$ ,  $CF_3SO_3$ ,  $CH_3SO_3$ ,  $HSO_4$ ,  $BF_4$ ,  $B(Phenyl)_4$ ,  $PF_6$ ,  $SbCl_6$ ,  $AsF_6$  or  $SbF_6$ ;  $X_8$  and  $X_9$  are the same or different and have the meaning of Z or  $A_2$ , or  $X_8$  has the meaning of Z or  $A_2$  and  $X_9$  stands for hydride.

49. (Previously Presented) A process according to claim 6, wherein  $R_1$  and/or  $R_2$  are, each independently,

pyrrolidinyl, piperidinyl, morpholinyl, tetrahydropyranyl, piperazinyl, pyrrolidinyl methyl, pyrrolidinyl ethyl, piperidinyl methyl, morpholinyl methyl, morpholinyl ethyl, tetrahydropyranyl methyl, tetrahydropyranyl ethyl, piperazinyl methyl or piperazinyl ethyl.

50-57. (Cancelled)